

**THERMAL TOLERANT AVICELASE FROM**  
**ACIDOTHERMUS CELLULOLYTICUS**

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**THERMAL TOLERANT AVICELASE FROM ACIDOTHERMUS CELLULOLYTICUS****Government Interests**

- 5 The United States Government has rights in this invention under Contract No. DE-AC36-99GO10337 between the United States Department of Energy and the National Renewable Energy Laboratory, a Division of the Midwest Research Institute.

**Field of the Invention**

- 10 The invention generally relates to a novel avicelase from *Acidothermus cellulolyticus*, AviIII. More specifically, the invention relates to purified and isolated AviIII polypeptides, nucleic acid molecules encoding the polypeptides, and processes for production and use of AviIII, as well as variants and derivatives thereof.

**Background of the Invention**

- 15 Plant biomass as a source of energy production can include agricultural and forestry products, associated by-products and waste, municipal solid waste, and industrial waste. In addition, over 50 million acres in the United States are currently available for biomass production, and there are a number of terrestrial and aquatic crops grown solely as a source for biomass (A Wiselogle, et al. Biomass feedstocks resources and composition. In CE Wyman, ed. Handbook on Bioethanol: Production and Utilization. Washington, DC: Taylor & Francis, 1996, pp 105-118). Biofuels produced from biomass include ethanol, methanol, biodiesel, and additives for reformulated gasoline. Biofuels are desirable because they add little, if any, net carbon dioxide to the atmosphere and because they greatly reduce ozone formation and carbon monoxide emissions as compared to the environmental output of conventional fuels. (P Bergeron. Environmental impacts of bioethanol. In CE Wyman, ed. Handbook on Bioethanol: Production and Utilization. Washington, DC: Taylor & Francis, 1996, pp 90-103).
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- 30 Plant biomass is the most abundant source of carbohydrate in the world due to the lignocellulosic materials composing the cell walls of all higher plants. Plant cell walls are divided into two sections, the primary and the secondary cell walls. The primary cell wall, which provides

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structure for expanding cells (and hence changes as the cell grows), is composed of three major polysaccharides and one group of glycoproteins. The predominant polysaccharide, and most abundant source of carbohydrates, is cellulose, while hemicellulose and pectin are also found in abundance. Cellulose is a linear beta-(1,4)-D-glucan and comprises 20% to 30% of the primary cell wall by weight. The secondary cell wall, which is produced after the cell has completed growing, also contains polysaccharides and is strengthened through polymeric lignin covalently cross-linked to hemicellulose.

Carbohydrates, and cellulose in particular can be converted to sugars by well-known methods including acid and enzymatic hydrolysis. Enzymatic hydrolysis of cellulose requires the processing of biomass to reduce size and facilitate subsequent handling. Mild acid treatment is then used to hydrolyze part or all of the hemicellulose content of the feedstock. Finally, cellulose is converted to ethanol through the concerted action of cellulases and saccharolytic fermentation (simultaneous saccharification fermentation (SSF)). The SSF process, using the yeast *Saccharomyces cerevisiae* for example, is often incomplete, as it does not utilize the entire sugar content of the plant biomass, namely the hemicellulose fraction.

The cost of producing ethanol from biomass can be divided into three areas of expenditure: pretreatment costs, fermentation costs, and other costs. Pretreatment costs include biomass milling, pretreatment reagents, equipment maintenance, power and water, and waste neutralization and disposal. The fermentation costs can include enzymes, nutrient supplements, yeast, maintenance and scale-up, and waste disposal. Other costs include biomass purchase, transportation and storage, plant labor, plant utilities, ethanol distillation, and administration (which may include technology-use licenses). One of the major expenses incurred in SSF is the cost of the enzymes, as about one kilogram of cellulase is required to fully digest 50 kilograms of cellulose. Economical production of cellulase is also compounded by factors such as the relatively slow growth rates of cellulase-producing organisms, levels of cellulase expression, and the tendency of enzyme-dependent processes to partially or completely inactivate enzymes due to conditions such as elevated temperature, acidity, proteolytic degradation, and solvent degradation.

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Enzymatic degradation of cellulose requires the coordinate action of at least three different types of cellulases. Such enzymes are given an Enzyme Commission (EC) designation according to the Nomenclature Committee of the International Union of Biochemistry and Molecular Biology (Eur. J. Biochem. 264: 607-609 and 610-650, 1999). Endo- beta-(1,4)-glucanases (EC 3.2.1.4) cleave the cellulose strand randomly along its length, thus generating new chain ends. Exo- beta-(1,4)-glucanases (EC 3.2.1.91) are processive enzymes and cleave cellobiosyl units (beta-(1,4)-glucose dimers) from free ends of cellulose strands. Lastly, beta-D-glucosidases (cellobiases: EC 3.2.1.21) hydrolyze cellobiose to glucose. All three of these general activities are required for efficient and complete hydrolysis of a polymer such as cellulose to a subunit, such as the simple sugar, glucose.

Highly thermostable enzymes have been isolated from the cellulolytic thermophile *Acidothermus cellulolyticus* gen. nov., sp. nov., a bacterium originally isolated from decaying wood in an acidic, thermal pool at Yellowstone National Park. A. Mohagheghi et al., (1986) Int. J. Systematic Bacteriology, 36(3): 435-443. One cellulase enzyme produced by this organism, the endoglucanase EI, is known to display maximal activity at 75 °C to 83°C. M.P. Tucker et al. (1989), Bio/Technology, 7(8): 817-820. E1 endoglucanase has been described in U.S. Patent 5,275,944. The *A. cellulolyticus* E1 endoglucanase is an active cellulase; in combination with the exocellulase CBH I from *Trichoderma reesei*, E1 gives a high level of saccharification and contributes to a degree of synergism. Baker JO et al. (1994), Appl. Biochem. Biotechnol., 45/46: 245-256. The gene coding EI catalytic and carbohydrate binding domains and linker peptide were described in U.S. Patent 5,536,655. E1 has also been expressed as a stable, active enzyme from a wide variety of hosts, including *E. coli*, *Streptomyces lividans*, *Pichia pastoris*, cotton, tobacco, and *Arabidopsis* (Dai Z, Hooker BS, Anderson DB, Thomas SR. Transgenic Res. 2000 Feb; 9(1):43-54).

The potential exists for the successful, commercial-scale expression of heterologous cellulases, and in particular novel cellulases with or without any one or more desirable properties such as thermal tolerance and resistance to acid inactivation, proteolytic inactivation, and solvent inactivation. Such expression can occur in filamentous fungi, bacteria, and other hosts.

There is a need within the art to generate alternative cellulase enzymes capable of assisting in the commercial-scale processing of cellulose to sugar for use in biofuel production. Against this backdrop the present invention has been developed. The potential exists for the successful, commercial-scale expression of heterologous cellulase polypeptides, and in particular novel cellulase polypeptides with or without any one or more desirable properties such as thermal tolerance, and partial or complete resistance to extreme pH inactivation, proteolytic inactivation, solvent inactivation, chaotropic agent inactivation, oxidizing agent inactivation, and detergent inactivation. Such expression can occur in fungi, bacteria, and other hosts.

### **Summary of the Invention**

The present invention provides AviIII, a novel member of the glycoside hydrolase (GH) family of enzymes, and in particular a thermal tolerant glycoside hydrolase useful in the degradation of cellulose. AviIII polypeptides of the invention include those having an amino acid sequence shown in SEQ ID NO:1, as well as polypeptides having substantial amino acid sequence identity to the amino acid sequence of SEQ ID NO:1 and useful fragments thereof, including, a catalytic domain having significant sequence similarity to the GH74 family, a first carbohydrate binding domain (type II) and a second carbohydrate binding domain (type III). See FIG 1.

The invention also provides a polynucleotide molecule encoding AviIII polypeptides and fragments of AviIII polypeptides, for example catalytic and carbohydrate binding domains. Polynucleotide molecules of the invention include those molecules having a nucleic acid sequence as shown in SEQ ID NO:2; those that hybridize to the nucleic acid sequence of SEQ ID NO:2 under high stringency conditions; and those having substantial nucleic acid identity with the nucleic acid sequence of SEQ ID NO:2.

The invention includes variants and derivatives of the AviIII polypeptides, including fusion proteins. For example, fusion proteins of the invention include AviIII polypeptide fused to a heterologous protein or peptide that confers a desired function. The heterologous protein or peptide can facilitate purification, oligomerization, stabilization, or secretion of the AviIII polypeptide, for example. As further examples, the heterologous polypeptide can provide enhanced activity, including catalytic or binding activity, for AviIII polypeptides, where the enhancement is either additive or synergistic. A fusion protein of an embodiment of the

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invention can be produced, for example, from an expression construct containing a polynucleotide molecule encoding AviIII polypeptide in frame with a polynucleotide molecule for the heterologous protein. Embodiments of the invention also comprise vectors, plasmids, expression systems, host cells, and the like, containing a AviIII polynucleotide molecule. Genetic engineering methods for the production of AviIII polypeptides of embodiments of the invention include expression of a polynucleotide molecule in cell free expression systems and in cellular hosts, according to known methods.

The invention further includes compositions containing a substantially purified AviIII polypeptide of the invention and a carrier. Such compositions are administered to a biomass containing cellulose for the reduction or degradation of the cellulose.

The invention also provides reagents, compositions, and methods that are useful for analysis of AviIII activity.

These and various other features as well as advantages which characterize the present invention will be apparent from a reading of the following detailed description and a review of the associated drawings.

The following Tables 4 and 5 includes sequences used in describing embodiments of the present invention. In Table 4, the abbreviations are as follows: CD, catalytic domain; CBD\_II, carbohydrate binding domain type II; CBD\_III, carbohydrate binding domain type III; and FN-III, fibronectin domain type III. When used herein, N\* indicates a string of unknown nucleic acid units, and X\* indicates a string of unknown amino acid units, for example about 50 or more. Table 4 includes approximate start and stop information for segments, and Table 5 includes amino acid sequence data for segments.

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Table 4. Nucleotide and polypeptide segments.

AvIII Segment	base BEGIN	base END	Length, bp	aa BEGIN No.	aa	aa END No.	aa	Length, aa	SEQ ID No. (amino acid)	SEQ ID No. (nucleotide)
Total length	1	about 3000	about 3 kb	1	M	about 1000	X*	about 1 kb	1	2
Signal (potential)	1	108	108	1	M	36	A	36		
CD (GH74)	109	2328	2220	37	A	776	G	740	3	
CBD_III (partial)	2575	about 3000	about 0.5 kb	859	V	about 1000	X*	about 154	4	
CBD_III (partial)	2575	2838	264	859	V	946	Q	88	5	

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Table 5. Gene/polypeptide segments with amino acid sequences.

SEQ ID No. (amino acid)	SEQ ID No. (nucleotide)	Aviii Segment	segment data	
			SEQ ID NO: 1 (see Table 1); SEQ ID NO: 2 (see Table 2)	
1	2	Total length		
		Signal (potential)	MRSRRLLVSLLAATAAFVAAALGVLPIAITASPAHA	
			<p>ATTQPYTWSNVAIGGGFVDGIVFNEGAPGILYVRTDIGMYRWDAAANGRWIPLLDWVGWNNWYNGV  VSIADPINTNKVWAAVGMVYNSWDPNMGAILRSSDQGWATQITPLPFKLGNMPPGRGMGERLAVDPN  NDNILYFGAPSGKGLWRSTDGATWSQMTNFPDVGTYIANPTDTTGYQSDIQGVVWVAFDKSSSSLGQ  ASKTIFVGVADPNPWFWRDGGATWQAVPGAPTGFIPHKGVFDPVNHVLYIATSNITGGPYDGSSEGDV  WKFSVTSGTWTRIISPVPSDTANDYFGYSGLTIDRQHPNTIMVATQISWWPDTIIFRSTDDGGATWTRI  WDWTSYPNRSRLRYVLDISAEPWLTFGVQPNPPVPSPKLGWMDAAMIDPFNSDRMLYGTGATLYATND  LTKWDSGGQIHIAPMVKGLEETAVNDLISPPSGAPLISALGDLGGFTHADVTAVPSTIFTSPVFTTGT  SVDYAEINPSIIVRAGSFDPSSQPNDRHVAFTDGGKNWFQSGEPGGVTTGGTVAASADGSRFVWAPG  DPGQPVVYAVGFGNSWAASQGVPANAQIRSDRVNPKTFYALSNGTFFYRSTDDGGVTFQPVAAGLPSSGA  VGVMFHAVPGKEGDLWLAASSGLYHSTNGGSSWSAITGVSSAVNVFGKSAPGSSYPAVFVVGTIGGV  TGAYRSDDCGTTWVLINDDQHQYGNWQQAITGDHANLRRVYIGTNGRGIYYGDI GGAPSG</p>	
3		CD (GH74)	<p>VSGGVKVQYKNND SAPGDNQIKPGLQVVNTGSSSSVDLSTVTVRYWFTDRDGGSTLVYNCDWAAIGCGN  IRASFGSVNPATPTADTYLQX*</p>	
4		CBD_III (partial)		
5		CBD_III (partial)	<p>VSGGVKVQYKNND SAPGDNQIKPGLQVVNTGSSSSVDLSTVTVRYWFTDRDGGSTLVYNCDWAAIGCGN  IRASFGSVNPATPTADTYLQ</p>	



### **Brief Description of the Drawings**

FIG. 1 is a schematic representation of the gene sequence and amino acid segment organization.

FIG 2 is a graphic representation of the glycoside hydrolase gene/protein families found in

5 various organisms.

### **Detailed Description**

#### **Definitions:**

The following definitions are provided to facilitate understanding of certain terms used frequently  
10 herein and are not meant to limit the scope of the present disclosure:

"Amino acid" refers to any of the twenty naturally occurring amino acids as well as any modified amino acid sequences. Modifications may include natural processes such as posttranslational processing, or may include chemical modifications which are known in the art. Modifications  
15 include but are not limited to: phosphorylation, ubiquitination, acetylation, amidation, glycosylation, covalent attachment of flavin, ADP-ribosylation, cross linking, iodination, methylation, and alike.

"Antibody" refers to a Y-shaped molecule having a pair of antigen binding sites, a hinge region  
20 and a constant region. Fragments of antibodies, for example an antigen binding fragment (Fab), chimeric antibodies, antibodies having a human constant region coupled to a murine antigen binding region, and fragments thereof, as well as other well known recombinant antibodies are included in the present invention.

25 "Antisense" refers to polynucleotide sequences that are complementary to target "sense" polynucleotide sequence.

"Binding activity" refers to any activity that can be assayed by characterizing the ability of a polypeptide to bind to a substrate. The substrate can be a polymer such as cellulose or can be  
30 complex molecule or aggregate of molecules where the entire moiety comprises at least some cellulose.

"Cellulase activity" refers to any activity that can be assayed by characterizing the enzymatic activity of a cellulase. For example, cellulase activity can be assayed by determining how much reducing sugar is produced during a fixed amount of time for a set amount of enzyme (see Irwin et al., (1998) *J. Bacteriology*, 1709-1714). Other assays are well known in the art and can be substituted.

"Complementary" or "complementarity" refers to the ability of a polynucleotide in a polynucleotide molecule to form a base pair with another polynucleotide in a second polynucleotide molecule. For example, the sequence A-G-T is complementary to the sequence T-C-A. Complementarity may be partial, in which only some of the polynucleotides match according to base pairing, or complete, where all the polynucleotides match according to base pairing.

"Expression" refers to transcription and translation occurring within a host cell. The level of expression of a DNA molecule in a host cell may be determined on the basis of either the amount of corresponding mRNA that is present within the cell or the amount of DNA molecule encoded protein produced by the host cell (Sambrook et al., 1989, *Molecular cloning: A Laboratory Manual*, 18.1-18.88).

"Fusion protein" refers to a first protein having attached a second, heterologous protein. Preferably, the heterologous protein is fused via recombinant DNA techniques, such that the first and second proteins are expressed in frame. The heterologous protein can confer a desired characteristic to the fusion protein, for example, a detection signal, enhanced stability or stabilization of the protein, facilitated oligomerization of the protein, or facilitated purification of the fusion protein. Examples of heterologous proteins useful in the fusion proteins of the invention include molecules having one or more catalytic domains of AviIII, one or more binding domains of AviIII, one or more catalytic domains of a glycoside hydrolase other than AviIII, one or more binding domains of a glycoside hydrolase other than AviIII, or any combination thereof. Further examples include immunoglobulin molecules and portions thereof, peptide tags such as histidine tag (6-His), leucine zipper, substrate targeting moieties, signal peptides, and the like. Fusion proteins are also meant to encompass variants and derivatives of AviIII polypeptides that

are generated by conventional site-directed mutagenesis and more modern techniques such as directed evolution, discussed infra.

"Genetically engineered" refers to any recombinant DNA or RNA method used to create a prokaryotic or eukaryotic host cell that expresses a protein at elevated levels, at lowered levels, or in a mutated form. In other words, the host cell has been transfected, transformed, or transduced with a recombinant polynucleotide molecule, and thereby been altered so as to cause the cell to alter expression of the desired protein. Methods and vectors for genetically engineering host cells are well known in the art; for example various techniques are illustrated in Current Protocols in Molecular Biology, Ausubel et al., eds. (Wiley & Sons, New York, 1988, and quarterly updates). Genetically engineering techniques include but are not limited to expression vectors, targeted homologous recombination and gene activation (see, for example, U.S. Patent No. 5,272,071 to Chappel) and trans activation by engineered transcription factors (see, for example, Segal et al., 1999, *Proc Natl Acad Sci USA* 96(6):2758-63).

"Glycoside hydrolase family" refers to a family of enzymes which hydrolyze the glycosidic bond between two or more carbohydrates or between a carbohydrate and a non-carbohydrate moiety (Henrissat B., (1991) *Biochem. J.*, 280:309-316). Identification of a putative glycoside hydrolase family member is made based on an amino acid sequence comparison and the finding of significant sequence similarity within the putative member's catalytic domain, as compared to the catalytic domains of known family members.

"Homology" refers to a degree of complementarity between polynucleotides, having significant effect on the efficiency and strength of hybridization between polynucleotide molecules. The term also can refer to a degree of similarity between polypeptides.

"Host cell" or "host cells" refers to cells expressing a heterologous polynucleotide molecule. Host cells of the present invention express polynucleotides encoding AviIII or a fragment thereof. Examples of suitable host cells useful in the present invention include, but are not limited to, prokaryotic and eukaryotic cells. Specific examples of such cells include bacteria of the genera *Escherichia*, *Bacillus*, and *Salmonella*, as well as members of the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*; fungi, particularly filamentous fungi such as *Trichoderma* and

*Aspergillus*, *Phanerochaete chrysosporium* and other white rot fungi; also other fungi including *Fusaria*, molds, and yeast including *Saccharomyces* sp., *Pichia* sp., and *Candida* sp. and the like; plants e.g. *Arabidopsis*, cotton, barley, tobacco, potato, and aquatic plants and the like; SF9 insect cells (Summers and Smith, 1987, *Texas Agriculture Experiment Station Bulletin*, 1555), and the like. Other specific examples include mammalian cells such as human embryonic kidney cells (293 cells), Chinese hamster ovary (CHO) cells (Puck et al., 1958, *Proc. Natl. Acad. Sci. USA* 60, 1275-1281), human cervical carcinoma cells (HELA) (ATCC CCL 2), human liver cells (Hep G2) (ATCC HB8065), human breast cancer cells (MCF-7) (ATCC HTB22), human colon carcinoma cells (DLD-1) (ATCC CCL 221), Daudi cells (ATCC CRL-213), murine myeloma cells such as P3/NSI/1-Ag4-1 (ATCC TIB-18), P3X63Ag8 (ATCC TIB-9), SP2/0-Ag14 (ATCC CRL-1581) and the like.

"Hybridization" refers to the pairing of complementary polynucleotides during an annealing period. The strength of hybridization between two polynucleotide molecules is impacted by the homology between the two molecules, stringency of the conditions involved, the melting temperature of the formed hybrid and the G:C ratio within the polynucleotides.

"Identity" refers to a comparison between pairs of nucleic acid or amino acid molecules. Methods for determining sequence identity are known. See, for example, computer programs commonly employed for this purpose, such as the Gap program (Wisconsin Sequence Analysis Package, Version 8 for Unix, Genetics Computer Group, University Research Park, Madison Wisconsin), that uses the algorithm of Smith and Waterman, 1981, *Adv. Appl. Math.*, 2: 482-489.

"Isolated" refers to a polynucleotide or polypeptide that has been separated from at least one contaminant (polynucleotide or polypeptide) with which it is normally associated. For example, an isolated polynucleotide or polypeptide is in a context or in a form that is different from that in which it is found in nature.

"Nucleic acid sequence" refers to the order or sequence of deoxyribonucleotides along a strand of deoxyribonucleic acid. The order of these deoxyribonucleotides determines the order of amino acids along a polypeptide chain. The deoxyribonucleotide sequence thus codes for the amino acid sequence.

“Polynucleotide” refers to a linear sequence of nucleotides. The nucleotides may be ribonucleotides, or deoxyribonucleotides, or a mixture of both. Examples of polynucleotides in the context of the present invention include single and double stranded DNA, single and double stranded RNA, and hybrid molecules having mixtures of single and double stranded DNA and RNA. The polynucleotides of the present invention may contain one or more modified nucleotides.

“Protein,” “peptide,” and “polypeptide” are used interchangeably to denote an amino acid polymer or a set of two or more interacting or bound amino acid polymers.

“Purify,” or “purified” refers to a target protein that is free from at least 5-10% of contaminating proteins. Purification of a protein from contaminating proteins can be accomplished using known techniques, including ammonium sulfate or ethanol precipitation, acid precipitation, heat precipitation, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography, size-exclusion chromatography, and lectin chromatography. Various protein purification techniques are illustrated in *Current Protocols in Molecular Biology*, Ausubel et al., eds. (Wiley & Sons, New York, 1988, and quarterly updates).

“Selectable marker” refers to a marker that identifies a cell as having undergone a recombinant DNA or RNA event. Selectable markers include, for example, genes that encode antimetabolite resistance such as the DHFR protein that confers resistance to methotrexate (Wigler et al, 1980, *Proc Natl Acad Sci USA* 77:3567; O'Hare et al., 1981, *Proc Natl Acad Sci USA*, 78:1527), the GPT protein that confers resistance to mycophenolic acid (Mulligan & Berg, 1981, *PNAS USA*, 78:2072), the neomycin resistance marker that confers resistance to the aminoglycoside G-418 (Calberre-Garapin et al., 1981, *J Mol Biol*, 150:1), the Hygro protein that confers resistance to hygromycin (Santerre et al., 1984, *Gene* 30:147), and the Zeocin™ resistance marker (Invitrogen). In addition, the herpes simplex virus thymidine kinase, hypoxanthine-guanine phosphoribosyltransferase and adenine phosphoribosyltransferase genes can be employed in tk<sup>-</sup>, hgprt<sup>-</sup> and aprt<sup>-</sup> cells, respectively.

"Stringency" refers to the conditions (temperature, ionic strength, solvents, etc) under which hybridization between polynucleotides occurs. A hybridization reaction conducted under high stringency conditions is one that will only occur between polynucleotide molecules that have a high degree of complementary base pairing (85% to 100% identity). Conditions for high stringency hybridization, for example, may include an overnight incubation at about 42°C for about 2.5 hours in 6 X SSC/0.1% SDS, followed by washing of the filters in 1.0 X SSC at 65°C, 0.1% SDS. A hybridization reaction conducted under moderate stringency conditions is one that will occur between polynucleotide molecules that have an intermediate degree of complementary base pairing (50% to 84% identity).

"Substrate targeting moiety" refers to any signal on a substrate, either naturally occurring or genetically engineered, used to target any AviIII polypeptide or fragment thereof to a substrate. Such targeting moieties include ligands that bind to a substrate structure. Examples of ligand/receptor pairs include carbohydrate binding domains and cellulose. Many such substrate-specific ligands are known and are useful in the present invention to target a AviIII polypeptide or fragment thereof to a substrate. A novel example is a AviIII carbohydrate binding domain that is used to tether other molecules to a cellulose-containing substrate such as a fabric.

"Thermal tolerant" refers to the property of withstanding partial or complete inactivation by heat and can also be described as thermal resistance or thermal stability. Although some variation exists in the literature, the following definitions can be considered typical for the optimum temperature range of stability and activity for enzymes: psychrophilic (below freezing to 10°C); mesophilic (10°C to 50°C); thermophilic (50°C to 75°C); and caldophilic (75°C to above boiling water temperature). The stability and catalytic activity of enzymes are linked characteristics, and the ways of measuring these properties vary considerably. For industrial enzymes, stability and activity are best measured under use conditions, often in the presence of substrate. Therefore, cellulases that must act on process streams of cellulose must be able to withstand exposure up to thermophilic or even caldophilic temperatures for digestion times in excess of several hours.

In encompassing a wide variety of potential applications for embodiments of the present invention, thermal tolerance refers to the ability to function in a temperature range of from about 15°C to about 100°C. A preferred range is from about 30°C to about 80°C. A highly preferred

range is from about 50°C to about 70°C. For example, a protein that can function at about 45°C is considered in the preferred range even though it may be susceptible to partial or complete inactivation at temperatures in a range above about 45°C and less than about 80°C. For polypeptides derived from organisms such as *Acidothermus*, the desirable property of thermal tolerance among is often accompanied by other desirable characteristics such as: resistance to extreme pH degradation, resistance to solvent degradation, resistance to proteolytic degradation, resistance to detergent degradation, resistance to oxidizing agent degradation, resistance to chaotropic agent degradation, and resistance to general degradation. Cowan DA in Danson MJ et al. (1992) The Archaeobacteria, Biochemistry and Biotechnology at 149-159, University Press, Cambridge, ISBN 1855780100. Here 'resistance' is intended to include any partial or complete level of residual activity. When a polypeptide is described as thermal tolerant it is understood that any one, more than one, or none of these other desirable properties can be present.

"Variant", as used herein, means a polynucleotide or polypeptide molecule that differs from a reference molecule. Variants can include nucleotide changes that result in amino acid substitutions, deletions, fusions, or truncations in the resulting variant polypeptide when compared to the reference polypeptide.

"Vector," "extra-chromosomal vector" or "expression vector" refers to a first polynucleotide molecule, usually double-stranded, which may have inserted into it a second polynucleotide molecule, for example a foreign or heterologous polynucleotide. The heterologous polynucleotide molecule may or may not be naturally found in the host cell, and may be, for example, one or more additional copy of the heterologous polynucleotide naturally present in the host genome. The vector is adapted for transporting the foreign polynucleotide molecule into a suitable host cell. Once in the host cell, the vector may be capable of integrating into the host cell chromosomes. The vector may optionally contain additional elements for selecting cells containing the integrated polynucleotide molecule as well as elements to promote transcription of mRNA from transfected DNA. Examples of vectors useful in the methods of the present invention include, but are not limited to, plasmids, bacteriophages, cosmids, retroviruses, and artificial chromosomes.

Within the application, unless otherwise stated, the techniques utilized may be found in any of several well-known references, such as: *Molecular Cloning: A Laboratory Manual* (Sambrook et al. (1989) *Molecular cloning: A Laboratory Manual*), *Gene Expression Technology* (Methods in Enzymology, Vol. 185, edited by D. Goeddel, 1991 Academic Press, San Diego, CA), "Guide to Protein Purification" in *Methods in Enzymology* (M.P. Deutscher, 3d., (1990) Academic Press, Inc.), *PCR Protocols: A Guide to Methods and Applications* (Innis et al. (1990) Academic Press, San Diego, CA), *Culture of Animal Cells: A Manual of Basic Technique*, 2<sup>nd</sup> ed. (R.I. Freshney (1987) Liss, Inc., New York, NY), and *Gene Transfer and Expression Protocols*, pp 109-128, ed. E.J. Murray, The Humana Press Inc., Clifton, N.J.).

### **O-Glycoside Hydrolases:**

Glycoside hydrolases are a large and diverse family of enzymes that hydrolyse the glycosidic bond between two carbohydrate moieties or between a carbohydrate and a non-carbohydrate moiety (See FIG. 2). Glycoside hydrolase enzymes are classified into glycoside hydrolase (GH) families based on significant amino acid similarities within their catalytic domains. Enzymes having related catalytic domains are grouped together within a family, (Henrissat et al., (1991) supra, and Henrissat et al. (1996), *Biochem. J.* 316:695-696), where the underlying classification provides a direct relationship between the GH domain amino acid sequence and how a GH domain will fold. This information ultimately provides a common mechanism for how the enzyme will hydrolyse the glycosidic bond within a substrate, *i.e.*, either by a retaining mechanism or inverting mechanism (Henrissat, B, (1991) supra).

Cellulases belong to the GH family of enzymes. Cellulases are produced by a variety of bacteria and fungi to degrade the  $\beta$ -1,4 glycosidic bond of cellulose and to so produce successively smaller fragments of cellulose and ultimately produce glucose. At present, cellulases are found within are at least 11 different GH families. Three different types of cellulase enzyme activities have been identified within these GH families: exo-acting cellulases which cleave successive disaccharide units from the non-reducing ends of a cellulose chain; endo-acting cellulases which randomly cleave successive disaccharide units within the cellulose chain; and  $\beta$ -glucosidases which cleave successive disaccharide units to glucose (J. W. Deacon, (1997) *Modern Mycology*, 3rd Ed., ISBN: 0-632-03077-1, 97-98).



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Many cellulases are characterized by having a multiple domain unit within their overall structure, a GH or catalytic domain is joined to a carbohydrate-binding domain (CBD) by a glycosylated linker peptide (Koivula et al., (1996) Protein Expression and Purification 8:391-400). As noted above, cellulases do not belong to any one family of GH domains, but rather have been identified within at least 11 different GH families to date. The CBD type domain increases the concentration of the enzyme on the substrate, in this case cellulose, and the linker peptide provides flexibility for both larger domains.

Conversion of cellulose to glucose is an essential step in the production of ethanol or other biofuels from biomass. Cellulases are an important component of this process, where approximately one kilogram of cellulase can digest fifty kilograms of cellulose. Within this process, thermostable cellulases have taken precedent, due to their ability to function at elevated temperatures and under other conditions including pH extremes, solvent presence, detergent presence, proteolysis, etc. (see Cowan DA (1992), supra).

Highly thermostable cellulase enzymes are secreted by the cellulolytic thermophile *Acidothermus cellulolyticus* (U.S. Patent Nos. 5,275,944 and 5,110,735). This bacterium was originally isolated from decaying wood in an acidic, thermal pool at Yellowstone National Park and deposited with the American Type Culture Collection (ATCC 43068) (Mohagheghi et al., (1986) *Int. J. System. Bacteriol.*, 36:435-443).

Recently, a thermostable cellulase, E1 endoglucanase, was identified and characterized from *Acidothermus cellulolyticus* (U.S. Patent No. 5,536,655). The E1 endoglucanase has maximal activity between 75 and 83°C and is active to a pH well below 5. Thermostable cellulase, and E1 endoglucanase, are useful in the conversion of biomass to biofuels, and in particular, are useful in the conversion of cellulose to glucose. Conversion of biomass to biofuel represents an extremely important alternative fuel source that is more environmentally friendly than conventional fuels, and provides a use, in some cases, for waste products.

### AviIII:

As described more fully in the Examples below, AviIII, a novel thermostable cellulase, has now been identified and characterized. The predicted amino acid sequence of AviIII (SEQ ID NO:1)

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has an organization characteristic of a cellulase enzyme. AviIII contains a carbohydrate binding domain - linker domain - catalytic domain -linker domain- fibronectin domain - linker domain - carbohydrate binding domain unit. In particular, AviIII includes a carbohydrate binding domain type III (CBDIII) (amino acids from about A35 to about A187), a GH74 catalytic domain (amino acids from about N231 to about P870), and a CBD<sub>II</sub> (amino acids from about G1021 to about S1121).

As discussed in more detail below (Example 2), significant amino acid similarity of AviIII to other cellulases identifies AviIII as a cellulase. In addition, the predicted amino acid sequence (SEQ ID NO: 1) indicates that a CBD type III domain is present as characterized by Tomme P. et al. (1995), in *Enzymatic Degradation of Insoluble Polysaccharides* (Saddler JN & Penner M, eds.), at 142-163, American Chemical Society, Washington. See also Tomme, P. & Claeyssens, M. (1989) *FEBS Lett.* 243, 239-243; Gilkes, N.R et al., (1988) *J.Biol.Chem.* 263, 10401-10407.

AviIII, as noted above, has a catalytic domain, identified as belonging to the GH74 family. The GH74 domain family includes a number of exoglucanases, for example, from *Cellulomonas fimi*, and exoglucanase E3 isolated from *Thermobifida fusca*. The GH74 members degrade substrate using an inverting mechanism. Being a member of the GH74 family of proteins identifies AviIII as potentially having cellulase activity.

AviIII is also a thermostable cellulase as it is produced by the thermophile *Acidothermus cellulolyticus*. As discussed, AviIII polypeptides can have other desirable characteristics (see Cowan DA (1992), supra). Like other members of the cellulase family, and in particular thermostable cellulases, AviIII polypeptides are useful in the conversion of biomass to biofuels and biofuel additives, and in particular, biofuels from cellulose. It is envisioned that AviIII polypeptides could be used for other purposes, for example in detergents, pulp and paper processing, food and feed processing, and in textile processes. AviIII polypeptides can be used alone or in combination with one or more other cellulases or glycoside hydrolases to perform the uses described herein or known within the relevant art, all of which are within the scope of the present disclosure.

**AviIII Polypeptides:**

AviIII polypeptides of the invention include isolated polypeptides having an amino acid sequence as shown below in Example 1; Table 1 and in SEQ ID NO:1, as well as variants and derivatives, including fragments, having substantial identity to the amino acid sequence of SEQ ID NO:1 and that retain any of the functional activities of AviIII. AviIII polypeptide activity can be determined, for example, by subjecting the variant, derivative, or fragment to a substrate binding assay or a cellulase activity assay such as those described in Irwin D et al., J. Bacteriology 180(7): 1709-1714 (April 1998).

**Table 1. AviIII amino acid sequence. (SEQ ID NO: 1)**

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MDRSENIRLTMRRLVSLAATASFVAAAALGVLP IAITASPAHAATTQ
PYTWSNVAIGGGGFVDGIVFNEGAPGILYVRTDIGGMYRWDAANGRWIPL
LDWVGWNNWGYNGVVSIAADPINTNKVWAAVGMYTNSWDPNDGAILRSSD
15 QGATWQITPLPFLGGMMPGRGMGERLAVDPNNDNILYFGAPSGKGLWRS
TDSGATWSQMTNFPDVGTYIANPTDTTGYQSDIQGVVWVAFDKSSSSLGQ
ASKTIFVGVADPNNPVFWSRDGGATWQAVPGAPTGFIPHKGVFDPVNHVL
YIATSNTGGPYDGSSGDVWKFSVTSGTWTRISPVPSTDANDYFGYSGLT
IDRQHPNTIMVATQISWWPDIIIFRSTDGGATWTRIWDWTSYPNRSRLRYV
20 LDISAEPWLTFGVQPNPPVSPKLGWMDEAMAIDPFNSDRMLYGTGATLY
ATNDLTKWDSGGQIHIAPMVKGLEETAVNDLISPPSGAPLISALGDLGGF
THADVTAVPSTIFTSPVFTTGTSDVYAEINPSIIIVRAGSFDPSQPNDRH
VAFSTDGKNWFQSGEPGGVTTGGTVAASADGSRFVWAPGDPGQPVVYAV
GFGNSWAASQGVPAQAIRSDRVNPKTFYALSNGTFYRSTDGGVTFQPVV
25 AGLPSSGAVGVFMFHAVPGEGDLWLAASSGLYHSTNGSSWSAITGVSSA
VNVGFGKSAPGSSYPVAVFVGTIGGVGTGAYRSDDCGTTWVLINDDQHGYG
NWGQAITGDHANLRRVYIGTNGRGIVYGDIGGAPSGSPSPSVSPSASPSL
SPSPSPSSSPSPSPSPSSSPSPSPSPSPSPSPSPSRSPSPSASPSPSPP
SPSSSPSSSPSPTPSSSPVSGGVKVQYKNDSAPGDNQIKPGLQVNTGS
30 SSVDLSTVTTRYWFTRDGGSSTLVYNCDWAAIGCGNIRASFGSVNPATPT
ADTYLQX*

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As listed and described in Tables 1 and 5, the isolated AviIII polypeptide includes an N-terminal hydrophobic region that functions as a signal peptide, having an amino acid sequence that begins with Met1 and extends to about A34; a carbohydrate binding domain having sequence similarity to such type III domains that begins with about A35 and extends to about A187, a catalytic domain having significant sequence similarity to a GH74 family domain that begins with about N231 and extends to about P870, a fibronectin type III domain that begins with about D901 and extends to about G985, a carbohydrate binding domain type II region that begins with about G1021 and extends to about S1121. Variants and derivatives of AviIII include, for example, AviIII polypeptides modified by covalent or aggregative conjugation with other chemical moieties, such as glycosyl groups, polyethylene glycol (PEG) groups, lipids, phosphate, acetyl groups, and the like.

The amino acid sequence of AviIII polypeptides of the invention is preferably at least about 60% identical, more preferably at least about 70% identical, or in some embodiments at least about 90% identical, to the AviIII amino acid sequence shown above in Table 1 and SEQ ID NO:1. The percentage identity, also termed homology (see definition above) can be readily determined, for example, by comparing the two polypeptide sequences using any of the computer programs commonly employed for this purpose, such as the Gap program (Wisconsin Sequence Analysis Package, Version 8 for Unix, Genetics Computer Group, University Research Park, Madison Wisconsin), which uses the algorithm of Smith and Waterman, 1981, *Adv. Appl. Math.* 2: 482-489.

Variants and derivatives of the AviIII polypeptide may further include, for example, fusion proteins formed of a AviIII polypeptide and a heterologous polypeptide. Preferred heterologous polypeptides include those that facilitate purification, oligomerization, stability, or secretion of the AviIII polypeptides.

AviIII polypeptide variants and derivatives, as used in the description of the invention, can contain conservatively substituted amino acids, meaning that one or more amino acid can be replaced by an amino acid that does not alter the secondary and/or tertiary structure of the polypeptide. Such substitutions can include the replacement of an amino acid, by a residue having similar physicochemical properties, such as substituting one aliphatic residue (Ile, Val, Leu, or Ala) for another, or substitutions between basic residues Lys and Arg, acidic residues Glu and Asp, amide residues Gln and Asn, hydroxyl residues Ser and Tyr, or aromatic residues Phe and Tyr. Phenotypically silent amino acid exchanges are described more fully in Bowie *et al.*, 1990, *Science* 247:1306-1310. In addition, functional AviIII polypeptide variants include those having amino acid substitutions, deletions, or additions to the amino acid sequence outside functional regions of the protein, for example, outside the catalytic and carbohydrate binding domains. These would include, for example, the various linker sequences that connect functional domains as defined herein.

The AviIII polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. The polypeptides may be recovered and purified from recombinant cell cultures by known methods, including, for example, ammonium sulfate or ethanol precipitation, anion or cation exchange chromatography, phosphocellulose chromatography,

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hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography, and lectin chromatography. Preferably, high performance liquid chromatography (HPLC) is employed for purification.

5 Another preferred form of AviIII polypeptides is that of recombinant polypeptides as expressed by suitable hosts. Furthermore, the hosts can simultaneously produce other cellulases such that a mixture is produced comprising a AviIII polypeptide and one or more other cellulases. Such a mixture can be effective in crude fermentation processing or other industrial processing.

10 AviIII polypeptides can be fused to heterologous polypeptides to facilitate purification. Many available heterologous peptides (peptide tags) allow selective binding of the fusion protein to a binding partner. Non-limiting examples of peptide tags include 6-His, thioredoxin, hemagglutinin, GST, and the OmpA signal sequence tag. A binding partner that recognizes and binds to the heterologous peptide can be any molecule or compound, including metal ions (for example, metal  
15 affinity columns), antibodies, antibody fragments, or any protein or peptide that preferentially binds the heterologous peptide to permit purification of the fusion protein.

AviIII polypeptides can be modified to facilitate formation of AviIII oligomers. For example, AviIII polypeptides can be fused to peptide moieties that promote oligomerization, such as leucine zippers and certain antibody fragment polypeptides, for example, Fc polypeptides. Techniques for  
20 preparing these fusion proteins are known, and are described, for example, in WO 99/31241 and in Cosman et.al., 2001 *Immunity* 14:123-133. Fusion to an Fc polypeptide offers the additional advantage of facilitating purification by affinity chromatography over Protein A or Protein G columns. Fusion to a leucine-zipper (LZ), for example, a repetitive heptad repeat, often with four or  
25 five leucine residues interspersed with other amino acids, is described in Landschultz et al., 1988, *Science*, 240:1759.

It is also envisioned that an expanded set of variants and derivatives of AviIII polynucleotides and/or polypeptides can be generated to select for useful molecules, where such expansion is  
30 achieved not only by conventional methods such as site-directed mutagenesis (SDM) but also by more modern techniques, either independently or in combination.

Site-directed-mutagenesis is considered an informational approach to protein engineering and can rely on high-resolution crystallographic structures of target proteins and some stratagem for specific amino acid changes (Van Den Burg, B.; Vriend, G.; Veltman, O.R.; Venema, G.; Eijssink, V.G.H. Proc. Nat. Acad. Sci. U.S. 1998, 95, 2056-2060). For example, modification of the amino acid

5 sequence of AviIII polypeptides can be accomplished as is known in the art, such as by introducing mutations at particular locations by oligonucleotide-directed mutagenesis (Walder et al., 1986, Gene, 42:133; Bauer et al., 1985, Gene 37:73; Craik, 1985, BioTechniques, 12-19; Smith et al., 1981, Genetic Engineering: Principles and Methods, Plenum Press; and U.S. Patent No. 4,518,584 and U.S. Patent No. 4,737,462). SDM technology can also employ the recent advent of

10 computational methods for identifying site-specific changes for a variety of protein engineering objectives (Hellenga, H.W. Nature Structural Biol. 1998, 5, 525-527).

The more modern techniques include, but are not limited to, non-informational mutagenesis techniques (referred to generically as "directed evolution"). Directed evolution, in conjunction with

15 high-throughput screening, allows testing of statistically meaningful variations in protein conformation (Arnold, F.H. Nature Biotechnol. 1998, 16, 617-618). Directed evolution technology can include diversification methods similar to that described by Cramer et al. (1998, Nature 391: 288-291), site-saturation mutagenesis, staggered extension process (StEP) (Zhao, H.; Giver, L.; Shao, Z.; Affholter, J.A.; Arnold, F.H. Nature Biotechnol. 1998, 16, 258-262), and DNA

20 synthesis/reassembly (U.S. Patent 5,965,408).

Fragments of the AviIII polypeptide can be used, for example, to generate specific anti-AviIII antibodies. Using known selection techniques, specific epitopes can be selected and used to generate monoclonal or polyclonal antibodies. Such antibodies have utility in the assay of AviIII

25 activity as well as in purifying recombinant AviIII polypeptides from genetically engineered host cells.

#### **AviIII Polynucleotides:**

The invention also provides polynucleotide molecules encoding the AviIII polypeptides discussed

30 above. AviIII polynucleotide molecules of the invention include polynucleotide molecules having the nucleic acid sequence shown in Table 2 and SEQ ID NO: 2, polynucleotide molecules that hybridize to the nucleic acid sequence of Table 2 and SEQ ID NO:2 under high stringency

hybridization conditions (for example, 42°, 2.5 hr., 6X SCC, 0.1%SDS); and polynucleotide molecules having substantial nucleic acid sequence identity with the nucleic acid sequence of Table 2 and SEQ ID NO:2, particularly with those nucleic acids encoding the catalytic domain, GH74 (from about amino acid A37 to about G776), the carbohydrate binding domain III (from about amino acid V859 to about at least Q946).

**Table 2. AviIII nucleotide sequence. (SEQ ID NO: 2)**

ATGGATCGTTCCGAGAACATCCGTCCTGACTATGAGATCACGACGATTGGTATCACTGCTCGCCGCCACTGCGTCGT  
 TCGCCGTGGCCGCCGCTCTGGGAGTTCTGCCCATCGCGATAACGGCTTCTCCTGCGCACGCGGCGACGACTCAGCC  
 GTACACCTGGAGCAACGTGGCGATCGGGGGCGCGGCTTTGTGCGACGGGATCGTCTTCAATGAAGGTGCACCGGGA  
 ATTCTGTACGTGCGGACGGACATCGGGGGGATGTATCGATGGGATGCCGCCAACGGGCGGTGGATCCCTCTTCTGG  
 ATTGGGTGGGATGGAACAATTGGGGGTACAACGGCGTCGTCAGCATTGCGGCAGACCCGATCAATACTAACAAGGT  
 ATGGGCCGCCGTCGGAATGTACACCAACAGCTGGGACCCAAACGACGGAGCGATTCTCCGCTCGTCTGATCAGGGC  
 GCAACGTGGCAAATAACGCCCCCTGCCGTTCAAGCTTGGCGGCAACATGCCCGGGCGTGGAATGGGCGAGCGGCTTG  
 CGGTGGATCCAAACAATGACAACATTTCTGTATTTCCGGCGCCCCGAGCGGCAAAGGGCTCTGGAGAAGCACAGATT  
 CGGCGCGACCTGGTCCCAGATGACGAACTTTCCGGACGTAGGCACGTACATTGCAAATCCCACGTACACGACCGGC  
 TATCAGAGCGATATTCAAGGCGTCGTCTGGGTGCGTTTCGACAAGTCTTCGTTCATCGCTCGGGCAAGCGAGTAAGA  
 CCATTTTTGTGGGCGTGGCGGATCCCAATAATCCGGTCTTCTGGAGCAGAGACGGCGGCGACGTGGCAGGCGGT  
 GCCGGGTGCGCCGACCGGCTTCATCCCGCACAAAGGCGCTTTTGACCCGGTCAACCACGTGCTCTATATTGCCACC  
 AGCAATACGGGTGGTCCGTATGACGGGAGCTCCGGCGACGTCTGGAAATTCTCGGTGACCTCCGGGACATGGACGC  
 GAATCAGCCCGGTACCTTCGACGGACACGGCCAACGACTACTTTGGTTACAGCGGCCTCACTATCGACCGCCAGCA  
 CCCGAACACGATAATGGTGGCAACCCAGATATCGTGGTGGCCGGACACCATAATCTTTCGGAGCACCGACGGCGGT  
 GCGACGTGGACGCGGATCTGGGATTTGGACGAGTTATCCCAATCGAAGCTTGCGATATGTGCTTGACATTTCCGCGG  
 AGCCTTGGCTGACCTTCGGCGTACAGCCGAATCCTCCCGTACCCAGTCCGAAGCTCGGCTGGATGGATGAAGCGAT  
 GGCAATCGATCCGTTCAACTCTGATCGGATGCTCTACGGAACAGGCGCGACGTTGTACGCAACAAATGATCTCAG  
 AAGTGGGACTCCGGCGGCGAGATTCAATCGCGCCGATGGTCAAAGGATTGGAGGAGACGGCGGTAAACGATCTCA  
 TCAGCCCGCGCTCTGGCGCCCCGCTCATCAGCGCTCTCGGAGACCTCGGCGGCTTCACCCACGCCGACGTTACTGC  
 CGTGCCATCGACGATCTTCACGTACCCGGTGTTCACGACCGGCACCGAGCGTCGACTATGCGGAATTGAATCCGTGC  
 ATCATCGTTTCGCGCTGGAAGTTTCGATCCATCGAGCCAACCGAACGACAGGCACGTGCGGTTCTCGACAGACGGCG  
 GCAAGAACTGGTTCCAAAGGCGAGCAACCTGGCGGGGTGACGACGGGCGGCACCGTCCCGCATCGGCCGACGGCTC  
 TCGTTTCGTCTGGGCTCCCGGCGATCCCGGTGAGCCTGTGGTGTACGAGTCGGATTTGGCAACTCCTGGGCTGCT  
 TCGCAAGGTGTTCCCGCAATGCCAGATCCGCTCAGACCGGGTGAATCCAAAGACTTTCTATGCCCTATCCAATG  
 GAACCTTCTATCGAAGCACGACGGCGGCGGTGACATTCCAACCGGTGCGGGCCGGTCTTCCGAGCAGCGGTGCCGT  
 CGGTGTCATGTTCCACGCGGTGCTTGGAAAAGAAAGCGATCTGTGGCTCGCTGCATCGAGCGGGCTTTACCACTCA  
 ACCAATGGCGGCAGCAGTTGGTCTGCAATCACCGGCGTATCCTCCGCGGTGAACGTGGGATTTGGTAAGTCTGCGC  
 CCGGGTCGTCATACCCAGCCGTCTTTGTGCTCGGCACGATCGGAGGCGTTACGGGGGCGTACCGCTCCGACGACTG  
 TGGGACGACCTGGTACTGATCAATGATGACCAGCACAATACGGAATTTGGGGACAAGCAATCACCGGTGACCAC  
 GCGAATTTACGGCGGGTGTACATAGGCACGAACGGCCGTGGAATTGTATACGGGGACATTGGTGGTGCGCCGTCCG  
 GATCGCCGTCTCCGTGCGGTGAGTCCGTGCGCTTCGCGGAGCCTGAGCCCGAGCCCGAGCCCGAGCAGCTCGCCATC  
 GCCGTGCGCGTCGCGGAGCTCGAGTCCATCCTCGTCGCGCTCTCCGTGCGCGTCACCATCGCCGAGTCCGTCTCGG  
 TCTCCGTACCATCGGCGTCCGCGAGCCCGTCTTCGTACCGAGCCCGTCTTCGTACCCGTCTTCGTGCGCGAGCC  
 CAACGCGCTCGTCGTGCGCGGTGTCGGGTGGGGTGAAGTGCAGTATAAGAATAATGATTCGGCGCCGGGTGATAA  
 TCAGATCAAGCCGGGTTTGAGGTGGTGAATACCGGTCGTGCTCGGTGGATTGTGCGACGGTGACGGTGCGGTAC  
 TGGTTACCCGGGATGGTGGCTCGTCGACACTGGTGTACAACGTGACTGGGCGGCGATCGGGTGTGGGAATATCC  
 GCGCCTCGTTCGGCTCGGTGAACCCGGCGACGCCGACGGCGGACACCTACCTGCAGN\*

The AviIII polynucleotide molecules of the invention are preferably isolated molecules encoding the AviIII polypeptide having an amino acid sequence as shown in Table 1 and SEQ ID NO:1, as well as derivatives, variants, and useful fragments of the AviIII polynucleotide. The AviIII polynucleotide

sequence can include deletions, substitutions, or additions to the nucleic acid sequence of Table 2 and SEQ ID NO: 1.

The AviIII polynucleotide molecule of the invention can be cDNA, chemically synthesized DNA, DNA amplified by PCR, RNA, or combinations thereof. Due to the degeneracy of the genetic code, two DNA sequences may differ and yet encode identical amino acid sequences. The present invention thus provides an isolated polynucleotide molecule having a AviIII nucleic acid sequence encoding AviIII polypeptide, where the nucleic acid sequence encodes a polypeptide having the complete amino acid sequences as shown in Table 1 and SEQ ID NO: 1, or variants, derivatives, and fragments thereof.

The AviIII polynucleotides of the invention have a nucleic acid sequence that is at least about 60% identical to the nucleic acid sequence shown in Table 2 and SEQ ID NO: 2, in some embodiments at least about 70% identical to the nucleic acid sequence shown in Table 2 and SEQ ID NO: 2, and in other embodiments at least about 90% identical to the nucleic acid sequence shown in Table 2 and SEQ ID NO: 2. Nucleic acid sequence identity is determined by known methods, for example by aligning two sequences in a software program such as the BLAST program (Altschul, S.F et al. (1990) J. Mol. Biol. 215:403-410, from the National Center for Biotechnology Information (<http://www.ncbi.nlm.nih.gov/BLAST/>)).

The AviIII polynucleotide molecules of the invention also include isolated polynucleotide molecules having a nucleic acid sequence that hybridizes under high stringency conditions (as defined above) to a the nucleic acid sequence shown in Table 2 and SEQ ID NO: 2. Hybridization of the polynucleotide is to about 15 contiguous nucleotides, or about 20 contiguous nucleotides, and in other embodiments about 30 contiguous nucleotides, and in still other embodiments about 100 contiguous nucleotides of the nucleic acid sequence shown in Table 2 and SEQ ID NO: 2.

Useful fragments of the AviIII-encoding polynucleotide molecules described herein, include probes and primers. Such probes and primers can be used, for example, in PCR methods to amplify and detect the presence of AviIII polynucleotides *in vitro*, as well as in Southern and Northern blots for analysis of AviIII. Cells expressing the AviIII polynucleotide molecules of the invention can also be identified by the use of such probes. Methods for the production and use of such primers and



probes are known. For PCR, 5' and 3' primers corresponding to a region at the termini of the AviIII polynucleotide molecule can be employed to isolate and amplify the AviIII polynucleotide using conventional techniques.

- 5 Other useful fragments of the AviIII polynucleotides include antisense or sense oligonucleotides comprising a single-stranded nucleic acid sequence capable of binding to a target AviIII mRNA (using a sense strand), or DNA (using an antisense strand) sequence.

#### **Vectors and Host Cells:**

- 10 The present invention also provides vectors containing the polynucleotide molecules of the invention, as well as host cells transformed with such vectors. Any of the polynucleotide molecules of the invention may be contained in a vector, which generally includes a selectable marker and an origin of replication, for propagation in a host. The vectors further include suitable transcriptional or translational regulatory sequences, such as those derived from a mammalian, microbial, viral, or insect genes, operably linked to the AviIII polynucleotide molecule. Examples of such regulatory sequences include transcriptional promoters, operators, or enhancers, mRNA ribosomal binding sites, and appropriate sequences which control transcription and translation. Nucleotide sequences are operably linked when the regulatory sequence functionally relates to the DNA encoding the target protein. Thus, a promoter nucleotide sequence is operably linked to a AviIII DNA sequence  
15 if the promoter nucleotide sequence directs the transcription of the AviIII sequence.  
20

- Selection of suitable vectors for the cloning of AviIII polynucleotide molecules encoding the target AviIII polypeptides of this invention will depend upon the host cell in which the vector will be transformed, and, where applicable, the host cell from which the target polypeptide is to be  
25 expressed. Suitable host cells for expression of AviIII polypeptides include prokaryotes, yeast, and higher eukaryotic cells, each of which is discussed below.

- The AviIII polypeptides to be expressed in such host cells may also be fusion proteins that include regions from heterologous proteins. As discussed above, such regions may be included to allow, for  
30 example, secretion, improved stability, or facilitated purification of the AviIII polypeptide. For example, a nucleic acid sequence encoding an appropriate signal peptide can be incorporated into an expression vector. A nucleic acid sequence encoding a signal peptide (secretory leader) may be

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fused in-frame to the AviIII sequence so that AviIII is translated as a fusion protein comprising the signal peptide. A signal peptide that is functional in the intended host cell promotes extracellular secretion of the AviIII polypeptide. Preferably, the signal sequence will be cleaved from the AviIII polypeptide upon secretion of AviIII from the cell. Non-limiting examples of signal sequences that can be used in practicing the invention include the yeast I-factor and the honeybee melatin leader in Sf9 insect cells.

Suitable host cells for expression of target polypeptides of the invention include prokaryotes, yeast, and higher eukaryotic cells. Suitable prokaryotic hosts to be used for the expression of these polypeptides include bacteria of the genera *Escherichia*, *Bacillus*, and *Salmonella*, as well as members of the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*. For expression in prokaryotic cells, for example, in *E. coli*, the polynucleotide molecule encoding AviIII polypeptide preferably includes an N-terminal methionine residue to facilitate expression of the recombinant polypeptide. The N-terminal Met may optionally be cleaved from the expressed polypeptide.

Expression vectors for use in prokaryotic hosts generally comprise one or more phenotypic selectable marker genes. Such genes encode, for example, a protein that confers antibiotic resistance or that supplies an auxotrophic requirement. A wide variety of such vectors are readily available from commercial sources. Examples include pSPORT vectors, pGEM vectors (Promega, Madison, WI), pPROEX vectors (LTI, Bethesda, MD), Bluescript vectors (Stratagene), and pQE vectors (Qiagen).

AviIII can also be expressed in yeast host cells from genera including *Saccharomyces*, *Pichia*, and *Kluveromyces*. Preferred yeast hosts are *S. cerevisiae* and *P. pastoris*. Yeast vectors will often contain an origin of replication sequence from a 2T yeast plasmid, an autonomously replicating sequence (ARS), a promoter region, sequences for polyadenylation, sequences for transcription termination, and a selectable marker gene. Vectors replicable in both yeast and *E. coli* (termed shuttle vectors) may also be used. In addition to the above-mentioned features of yeast vectors, a shuttle vector will also include sequences for replication and selection in *E. coli*. Direct secretion of the target polypeptides expressed in yeast hosts may be accomplished by the inclusion of nucleotide sequence encoding the yeast I-factor leader sequence at the 5' end of the AviIII-encoding nucleotide sequence.

Insect host cell culture systems can also be used for the expression of AviIII polypeptides. The target polypeptides of the invention are preferably expressed using a baculovirus expression system, as described, for example, in the review by Luckow and Summers, 1988 *Bio/Technology* 6:47.

5

The choice of a suitable expression vector for expression of AviIII polypeptides of the invention will depend upon the host cell to be used. Examples of suitable expression vectors for *E. coli* include pET, pUC, and similar vectors as is known in the art. Preferred vectors for expression of the AviIII polypeptides include the shuttle plasmid pIJ702 for *Streptomyces lividans*,  
10 pGAPZalpha-A, B, C and pPICZalpha-A, B, C (Invitrogen) for *Pichia pastoris*, and pFE-1 and pFE-2 for filamentous fungi and similar vectors as is known in the art.

Modification of a AviIII polynucleotide molecule to facilitate insertion into a particular vector (for example, by modifying restriction sites), ease of use in a particular expression system or host  
15 (for example, using preferred host codons), and the like, are known and are contemplated for use in the invention. Genetic engineering methods for the production of AviIII polypeptides include the expression of the polynucleotide molecules in cell free expression systems, in cellular hosts, in tissues, and in animal models, according to known methods.

## 20 **Compositions**

The invention provides compositions containing a substantially purified AviIII polypeptide of the invention and an acceptable carrier. Such compositions are administered to biomass, for example, to degrade the cellulose in the biomass into simpler carbohydrate units and ultimately, to sugars. These released sugars from the cellulose are converted into ethanol by any number of  
25 different catalysts. Such compositions may also be included in detergents for removal, for example, of cellulose containing stains within fabrics, or compositions used in the pulp and paper industry, to address conditions associated with cellulose content. Compositions of the present invention can be used in stonewashing jeans such as is well known in the art. Compositions can be used in the biopolishing of cellulosic fabrics, such as cotton, linen, rayon and Lyocell.

30

The invention provides pharmaceutical compositions containing a substantially purified AviIII polypeptide of the invention and if necessary a pharmaceutically acceptable carrier. Such

pharmaceutical compositions are administered to cells, tissues, or patients, for example, to aid in delivery or targeting of other pharmaceutical compositions. For example, AviIII polypeptides may be used where carbohydrate-mediated liposomal interactions are involved with target cells. Vyas SP et al. (2001), J. Pharmacy & Pharmaceutical Sciences May-Aug 4(2): 138-58.

The invention also provides reagents, compositions, and methods that are useful for analysis of AviIII activity and for the analysis of cellulose breakdown.

Compositions of the present invention may also include other known cellulases, and preferably, other known thermal tolerant cellulases for enhanced treatment of cellulose.

### Antibodies

The polypeptides of the present invention, in whole or in part, may be used to raise polyclonal and monoclonal antibodies that are useful in purifying AviIII, or detecting AviIII polypeptide expression, as well as a reagent tool for characterizing the molecular actions of the AviIII polypeptide. Preferably, a peptide containing a unique epitope of the AviIII polypeptide is used in preparation of antibodies, using conventional techniques. Methods for the selection of peptide epitopes and production of antibodies are known. See, for example, *Antibodies: A Laboratory Manual*, Harlow and Land (eds.), 1988 Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y.; *Monoclonal Antibodies, Hybridomas: A New Dimension in Biological Analyses*, Kennet et al. (eds.), 1980 Plenum Press, New York.

### Assays

Agents that modify, for example, increase or decrease, AviIII hydrolysis or degradation of cellulose can be identified, for example, by assay of AviIII cellulase activity and/or analysis of AviIII binding to a cellulose substrate. Incubation of cellulose in the presence of AviIII and in the presence or absence of a test agent and correlation of cellulase activity or carbohydrate binding permits screening of such agents. For example, cellulase activity and binding assays may be performed in a manner similar to those described in Irwin et al., J. Bacteriology 180(7): 1709-1714 (April 1998).

The AviIII stimulated activity is determined in the presence and absence of a test agent and then compared. A lower AviIII activated test activity in the presence of the test agent, than in the absence of the test agent, indicates that the test agent has decreased the activity of the AviIII. A higher AviIII activated test activity in the presence of the test agent than in the absence of the test agent indicates that the test agent has increased the activity of the AviIII. Stimulators and inhibitors of AviIII may be used to augment, inhibit, or modify AviIII mediated activity, and therefore may have potential industrial uses as well as potential use in the further elucidation of AviIII's molecular actions.

### 10 **Therapeutic Applications**

The AviIII polypeptides of the invention are effective in adding in delivery or targeting of other pharmaceutical compositions within a host. For example, AviIII polypeptides may be used where carbohydrate-mediated liposomal interactions are involved with target cells. Vyas SP et al. (2001), *J. Pharm Pharm Sci* May-Aug 4(2): 138-58.

AviIII polynucleotides and polypeptides, including vectors expressing AviIII, of the invention can be formulated as pharmaceutical compositions and administered to a host, preferably mammalian host, including a human patient, in a variety of forms adapted to the chosen route of administration. The compounds are preferably administered in combination with a pharmaceutically acceptable carrier, and may be combined with or conjugated to specific delivery agents, including targeting antibodies and/or cytokines.

AviIII can be administered by known techniques, such as orally, parentally (including subcutaneous injection, intravenous, intramuscular, intrasternal or infusion techniques), by inhalation spray, topically, by absorption through a mucous membrane, or rectally, in dosage unit formulations containing conventional non-toxic pharmaceutically acceptable carriers, adjuvants or vehicles. Pharmaceutical compositions of the invention can be in the form of suspensions or tablets suitable for oral administration, nasal sprays, creams, sterile injectable preparations, such as sterile injectable aqueous or oleagenous suspensions or suppositories.

For oral administration as a suspension, the compositions can be prepared according to techniques well-known in the art of pharmaceutical formulation. The compositions can contain

microcrystalline cellulose for imparting bulk, alginic acid or sodium alginate as a suspending agent, methylcellulose as a viscosity enhancer, and sweeteners or flavoring agents. As immediate release tablets, the compositions can contain microcrystalline cellulose, starch, magnesium stearate and lactose or other excipients, binders, extenders, disintegrants, diluents and lubricants known in the art.

For administration by inhalation or aerosol, the compositions can be prepared according to techniques well-known in the art of pharmaceutical formulation. The compositions can be prepared as solutions in saline, using benzyl alcohol or other suitable preservatives, absorption promoters to enhance bioavailability, fluorocarbons or other solubilizing or dispersing agents known in the art.

For administration as injectable solutions or suspensions, the compositions can be formulated according to techniques well-known in the art, using suitable dispersing or wetting and suspending agents, such as sterile oils, including synthetic mono- or diglycerides, and fatty acids, including oleic acid.

For rectal administration as suppositories, the compositions can be prepared by mixing with a suitable non-irritating excipient, such as cocoa butter, synthetic glyceride esters or polyethylene glycols, which are solid at ambient temperatures, but liquefy or dissolve in the rectal cavity to release the drug.

Preferred administration routes include orally, parenterally, as well as intravenous, intramuscular or subcutaneous routes. More preferably, the compounds of the present invention are administered parenterally, i.e., intravenously or intraperitoneally, by infusion or injection.

Solutions or suspensions of the compounds can be prepared in water, isotonic saline (PBS) and optionally mixed with a nontoxic surfactant. Dispersions may also be prepared in glycerol, liquid polyethylene, glycols, DNA, vegetable oils, triacetin and mixtures thereof. Under ordinary conditions of storage and use, these preparations may contain a preservative to prevent the growth of microorganisms.

The pharmaceutical dosage form suitable for injection or infusion use can include sterile, aqueous solutions or dispersions or sterile powders comprising an active ingredient which are adapted for the extemporaneous preparation of sterile injectable or infusible solutions or dispersions. In all cases, the ultimate dosage form should be sterile, fluid and stable under the conditions of manufacture and storage. The liquid carrier or vehicle can be a solvent or liquid dispersion medium comprising, for example, water, ethanol, a polyol such as glycerol, propylene glycol, or liquid polyethylene glycols and the like, vegetable oils, nontoxic glyceryl esters, and suitable mixtures thereof. The proper fluidity can be maintained, for example, by the formation of liposomes, by the maintenance of the required particle size, in the case of dispersion, or by the use of nontoxic surfactants. The prevention of the action of microorganisms can be accomplished by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal, and the like. In many cases, it will be desirable to include isotonic agents, for example, sugars, buffers, or sodium chloride. Prolonged absorption of the injectable compositions can be brought about by the inclusion in the composition of agents delaying absorption--for example, aluminum monostearate hydrogels and gelatin.

Sterile injectable solutions are prepared by incorporating the compounds in the required amount in the appropriate solvent with various other ingredients as enumerated above and, as required, followed by filter sterilization. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying and freeze-drying techniques, which yield a powder of the active ingredient plus any additional desired ingredient present in the previously sterile-filtered solutions.

### **Industrial Applications**

The AviIII polypeptides of the invention are effective cellulases. In the methods of the invention, the cellulose degrading effects of AviIII are achieved by treating biomass at a ratio of about 1 to about 50, or about 1:40, 1:35, 1:30, 1:25, 1:20 or even about 1: 70 in some preparations of the AVIII of AviIII:biomass. AviIII may be used under extreme conditions, for example, elevated temperatures and acidic pH. Treated biomass is degraded into simpler forms of carbohydrates, and in some cases glucose, which is then used in the formation of ethanol or other industrial chemicals, as is known in the art. Other methods are envisioned to be within the scope of the present invention, including methods for treating fabrics to remove cellulose-containing stains

and other methods already discussed. AviIII polypeptides can be used in any known application currently utilizing a cellulase, all of which are within the scope of the present invention.

Having generally described the invention, the same will be more readily understood by reference to the following examples, which are provided by way of illustration and are not intended as limiting.

## EXAMPLES

### 10 Example 1: Molecular Cloning of AviIII

Genomic DNA was isolated from *Acidothermus cellulolyticus* and purified by banding on cesium chloride gradients. Genomic DNA was partially digested with Sau 3A and separated on agarose gels. DNA fragments in the range of 9-20 kilobase pairs were isolated from the gels. This purified Sau 3A digested genomic DNA was ligated into the Bam H1 acceptor site of purified EMBL3 lambda phage arms (Clontech, San Diego, Calif.). Phage DNA was packaged according to the manufacturer's specification and plated with *E. Coli* LE392 in top agar which contained the soluble cellulose analog, carboxymethylcellulose (CMC). The plates were incubated overnight (12-24 hours) to allow transfection, bacterial growth, and plaque formation. Plates were stained with Congo Red followed by destaining with 1 M NaCl. Lambda plaques harboring endoglucanase clones showed up as unstained plaques on a red background.

Lambda clones which screened positive on CMC-Congo Red plates were purified by successive rounds of picking, plating and screening. Individual phage isolates were named SL-1, SL-2, SL-3, and SL-4. Subsequent subcloning efforts employed the SL-3 clone which contained an approximately 14.2 kilobase fragment of *Acidothermus cellulolyticus* genomic DNA.

Template DNA was constructed using a 9 kilobase Bam H1 fragment obtained from the 14.2 kilobase lambda clone SL-3 prepared from *Acidothermus cellulolyticus* genomic DNA. The 9 kilobase Bam H1 fragment from SL-3 was subcloned into pDR540 to generate a plasmid NREL501. NREL501 was sequenced by the primer walking method as is known in the art. NREL501 was then subcloned into pUC19 using restriction enzymes Pst I and Eco RI and transformed into *E. coli* XL1-blue (Stratagene) for the production of template DNA for



sequencing. Each subclone was sequenced from both the forward and reverse directions. DNA for sequencing was prepared from an overnight growth in 500 mL LB broth using a megaprep DNA purification kit from Promega. The templated DNA was PEG precipitated and suspended in de-ionized water and adjusted to a final concentration of 0.25 milligrams/mL.

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Custom primers were designed by reading upstream known sequence and selecting segments of an appropriate length to function, as is well known in the art. Primers for cycle sequencing were synthesized at the Macromolecular Resources Facility located at Colorado State University in Fort Collins , Colorado. Typically the sequencing primers were 26 to 30 nucleotides in length, but were sometimes longer or shorter to accommodate a melting temperature appropriate for cycle sequencing. The sequencing primers were diluted in de-ionized water, the concentration measured using UV absorbance at 260 nm, and then adjusted to a final concentration of 5 pmol/microL.

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15 Templates and sequencing primers were shipped to the Iowa State University DNA Sequencing Facility at Ames, Iowa for sequencing using standard chemistries for cycle sequencing. In some cases, regions of the template that sequenced poorly using the standard protocols and dye terminators were repeated with the addition of 2 microL DMSO and by using nucleotides optimized for the sequencing of high GC content DNA. An inverse PCR technique known in the art was applied to continue sequencing the genomic DNA, and a primer walking method was used to sequence the large PCR products. Each PCR fragment was sequenced from both strands, using high fidelity commercial DNA polymerase.

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Sequencing data from primer walking and subclones were assembled together to verify that all SL-3 regions had been sequenced from both strands. An open reading frame (ORF) was found in the 9 kilobase Bam H1 fragment, C-terminal of E1 (U.S. Patent 5,536,655), termed AviIII. An ORF of 3366 bp [SEQ ID NO:2] and deduced amino acid sequence [SEQ ID NO:1] are shown in Tables 1 and 2. The amino acid sequence predicted by SEQ ID NO:1 was determined to have significant homology to known cellulases, as is shown below in Example 2 and Table 3.

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Sequence alignments and comparisons of the amino acid sequences of the *Acidothermus*

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Multialignment of related Glycoside Hydrolase Family 74 catalytic domain  
GH74\_Ace: *Acidothermus cellulolyticus* AviIII catalytic domain GH74  
AviIII\_Aac: *Aspergillus aculeatus* Avicelase III (endoglucanase). GeneBank Acc. # BAA29031

From the putative locations of the domains in the AviIII cellulase sequence given above and in comparable cloned cellulase sequences from other species, one can separate individual domains and combine them with one or more domains from different sequences. The significant similarity between cellulase genes permit one by recombinant techniques to arrange one or more domains

from the *Acidothermus cellulolyticus* AviIII cellulase gene with one or more domains from a cellulase gene from one or more other microorganisms. Other representative endoglucanase genes include *Bacillus polymyxa* beta-(1,4) endoglucanase (Baird et al, Journal of Bacteriology, 172: 1576-86 (1992)) and *Xanthomonas campestris* beta-(1,4)-endoglucanase A (Gough et al, 5 Gene 89:53-59 (1990)). The result of the fusion of any two or more domains will, upon expression, be a hybrid polypeptide. Such hybrid polypeptides can have one or more catalytic or binding domains. For ease of manipulation, recombinant techniques may be employed such as the addition of restriction enzyme sites by site-specific mutagenesis. If one is not using one domain of a particular gene, any number of any type of change including complete deletion may be made 10 in the unused domain for convenience of manipulation.

It is understood for purposes of this disclosure, that various changes and modifications may be made to the invention that are well within the scope of the invention. Numerous other changes may be made which will readily suggest themselves to those skilled in the art and which are 15 encompassed in the spirit of the invention disclosed herein and as defined in the appended claims.

This specification contains numerous citations to references such as patents, patent applications, and publications. Each is hereby incorporated by reference for all purposes.